

PCTWORLD INTELLECTUAL PI
Internation

INTERNATIONAL APPLICATION PUBLISHED UN

WO 9606930A1

(51) International Patent Classification ⁶ : C12N 9/02, C11D 3/386 // C12N 9/02, C12R 1:645		A1	(11) International Publication Number: WO 96/06930
			(43) International Publication Date: 7 March 1996 (07.03.96)
(21) International Application Number: PCT/DK95/00344			(81) Designated States: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TT, UA, UG, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, MW, SD, SZ, UG).
(22) International Filing Date: 25 August 1995 (25.08.95)			
(30) Priority Data: 0982/94 26 August 1994 (26.08.94) DK			
(71) Applicant (for all designated States except US): NOVO NORDISK A/S [DK/DK]; Novo Allé, DK-2880 Bagsværd (DK).			
(72) Inventors; and (75) Inventors/Applicants (for US only): DAMHUS, Ture [DK/DK]; Novo Nordisk A/S, Novo Allé, DK-2880 Bagsværd (DK). SCHNEIDER, Palle [DK/DK]; Novo Nordisk A/S, Novo Allé, DK-2880 Bagsværd (DK). BECH, Lisbeth [DK/DK]; Novo Nordisk A/S, Novo Allé, DK-2880 Bagsværd (DK). HEINZKILL, Marion [DE/DE]; Universität Kaiserslautern, Paul-Ehrlich-Strasse 23, D-67663 Kaiserslautern (DE).			
(74) Common Representative: NOVO NORDISK A/S; Corporate Patents, Novo Allé, DK-2880 Bagsværd (DK).			Published With international search report.

(54) Title: COPRINACEAE LACCASES**(57) Abstract**

The present invention relates to a method of oxidizing a substrate in a solution with a pH at or above 7, comprising contacting the substrate with an effective amount of a laccase obtainable from *Coprinaceae*.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgyzstan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

COPRINACEAE LACCASES

FIELD OF INVENTION

The present invention relates to a method of oxidizing a substrate in an alkaline solution with a laccase. The invention also relates to a detergent additive and to a detergent composition.

BACKGROUND OF THE INVENTION

Laccases are enzymes that catalyze the oxidation of a substrate with dioxygen, reducing dioxygen to water. Such enzymes are known from microbial, plant and animal origins.

It has earlier been found that coloured substances leached from dyed fabrics could be bleached in solution by means of peroxidases or suitable oxidases. The use of peroxidases or oxidases for inhibiting dye transfer in this way is described in WO 91/05839.

Certain oxidizable substrates, e.g., metal ions and phenolic compounds such as 7-hydroxycoumarin, vanillin, and p-hydroxybenzenesulfonate, have been described as accelerators or enhancing agents able to enhance bleaching reactions (cf. e.g. WO 92/18683, WO 92/18687, and Kato M and Shimizu S, Plant Cell Physiol. 1985 26 (7), pp. 1291-1301 (cf. Table 1 in particular)). In WO 94/12621 other types of enhancing agents are disclosed, e.g., phenothiazines and phenoxazines.

It is the object of this invention to provide a laccase system with bleaching performance at high pH.

SUMMARY OF THE INVENTION

We have unexpectedly found that laccases obtainable from Coprinaceae show excellent performance in bleaching assays at high pH.

Accordingly, the invention provides a method of oxidizing a substrate in a solution with a pH at or above 7, comprising contacting the substrate with an effective amount of

a laccase obtainable from Coprinaceae. The invention also provides use of these laccases in bleaching compositions and detergent compositions.

BRIEF DESCRIPTION OF DRAWINGS

5 The present invention is further illustrated by reference to Fig. 1 which shows the bleaching of gradually added Acid Blue 45 in phosphate/borate buffer pH 10 at 35°C; (I): Only dye addition; (II): Dye addition in the presence of Laccase; (III): Dye addition in the presence of Laccase + 10-ethylphenothiazine-4-carboxylic acid; (IV): Dye addition in the
10 presence of Laccase + phenoxazine-10-propionic acid; the experiment conducted as described in Example 4.

DETAILED DISCLOSURE OF THE INVENTION

 The present invention provides a method of oxidizing
15 a substrate in a solution with a pH at or above 7, preferably in a solution with a pH at or above 8, even more preferably in a solution with a pH at or above 9, in particular in a solution with a pH around 10, comprising contacting said substrate with an effective amount of a laccase obtainable from Coprinaceae,
20 optionally in the presence of an enhancing agent.

 Laccases (EC 1.10.3.2) are oxidoreductases that function with molecular oxygen as electron acceptor. Molecular oxygen from the atmosphere will usually be present in sufficient quantity, so normally it is not necessary to add extra
25 oxygen to the process medium.

Coprinaceae Laccases

 According to the invention useful laccases that function at a high pH are obtainable from laccase-producing strains of the family Coprinaceae.

30 The family Coprinaceae comprises the following genera: Coprinus, Podaxis, Montagnea, Macrometrula,

Psathyrella, Panaeolina, Panaeolus, Copelandia, Anellaria, Limnoperdon, Panaeolopsis and Polyplocium.

Preferably, the laccase employed in the method of the present invention is obtainable from Coprinus, Panaeolus or Psathyrella, in particular Coprinus cinereus, Coprinus comatus, Coprinus friesii, Coprinus plicatilis, or Psathyrella condolleana, or Panaeolus papilionaceus; most preferred strain is Coprinus cinereus (IFO30116). The most preferred strain is freely available to the public from Institute of Fermentation, Osaka (IFO) under the indicated deposit number.

A strain representative of Coprinus comatus has been deposited according to the Budapest Treaty on the International Recognition of the Deposits of Microorganisms for the Purpose of Patent Procedures, on 16 August 1995, at Centraalbureau voor Schimmelcultures, Oosterstraat 1, Postbus 273, NL-3740 AG Baarn, Netherlands, under Accession No. CBS 631.95.

A strain representative of Coprinus friesii has been deposited according to the Budapest Treaty on the International Recognition of the Deposits of Microorganisms for the Purpose of Patent Procedures, on 16 August 1995, at Centraalbureau voor Schimmelcultures, Oosterstraat 1, Postbus 273, NL-3740 AG Baarn, Netherlands, under Accession No. CBS 629.95.

A strain representative of Coprinus plicatilis has been deposited according to the Budapest Treaty on the International Recognition of the Deposits of Microorganisms for the Purpose of Patent Procedures, on 16 August 1995, at Centraalbureau voor Schimmelcultures, Oosterstraat 1, Postbus 273, NL-3740 AG Baarn, Netherlands, under Accession No. CBS 627.95.

A strain representative of Psathyrella condolleana has been deposited according to the Budapest Treaty on the International Recognition of the Deposits of Microorganisms for the Purpose of Patent Procedures, on 16 August 1995, at Centraalbureau voor Schimmelcultures, Oosterstraat 1, Postbus 273, NL-3740 AG Baarn, Netherlands, under Accession No. CBS 628.95.

A strain representative of Panaeolus papilionaceus has been deposited according to the Budapest Treaty on the

International Recognition of the Deposits of Microorganisms for the Purpose of Patent Procedures, on 16 August 1995, at Centraalbureau voor Schimmelcultures, Oosterstraat 1, Postbus 273, NL-3740 AG Baarn, Netherlands, under Accession No. CBS 5 630.95.

The laccase according to the invention may furthermore be one which is producible by a method comprising cultivating a host cell transformed with a recombinant DNA vector which carries a DNA sequence encoding said laccase as well as 10 DNA sequences encoding functions permitting the expression of the DNA sequence encoding the laccase, in a culture medium under conditions permitting the expression of the laccase enzyme, and recovering the laccase from the culture.

Determination of Laccase Activity (LACU)

15 Laccase activity is determined by oxidizing syringaldazin under aerobic conditions. The violet colour produced is photometered at 530 nm. The analytical conditions are 19 μ M syringaldazin, 23.2 mM acetate buffer, pH 5.5, 30°C, 1 min reaction time.

20 1 laccase unit (LACU) is the amount of enzyme that catalyses the conversion of 1.0 μ mole syringaldazin per minute under these conditions.

Immunochemical Properties

The immunochemical properties can be determined 25 immunologically by cross-reaction identity tests. The identity tests can be performed by the well-known Ouchterlony double immunodiffusion procedure or by tandem crossed immunoelectrophoresis according to I. M. Roitt; Immunology, Gower Medical Publishing (1985) and N. H. Axelsen; Handbook of Immunoprecipitation-in-Gel Techniques; Blackwell Scientific Publications 30 (1983), Chapters 5 and 14.

According to the invention laccases displaying immunochemical cross-reactivity with an antibody raised against

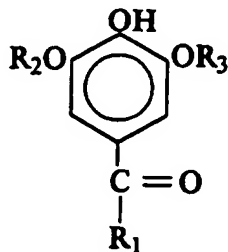
a laccase obtainable from Coprinus cinereus IFO30116 are also included.

Enhancing Agents

In oxidizing a substrate at high pH the laccases of the invention show excellent results. If additionally an enhancing agent is added, the result may be improved.

According to the invention an enhancing agent is any compound that enhances the oxidation. The enhancing agent will typically be an oxidizable compound, e.g., a metal ion or a phenolic compound such as 7-hydroxycoumarin, vanillin, or p-hydroxybenzenesulfonate, (for reference see WO 92/18683, WO 92/18687, and Kato M and Shimizu S, Plant Cell Physiol. 1985 26 (7), pp. 1291-1301 (cf. Table 1 in particular)).

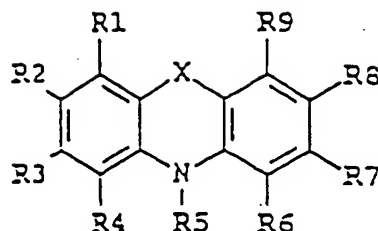
Particularly useful enhancing agents may be described by the following formula:



where R_1 is H, OH, $\text{C}_n\text{H}_{2n+1}$ or $\text{OC}_n\text{H}_{2n+1}$, in which n is an integer of from 1 to 10; and R_2 and R_3 are the same or different and selected from $\text{C}_m\text{H}_{2m+1}$, in which m is an integer of from 1 to 10. R_1 , R_2 and R_3 may also contain double bonds or cyclic groups.

In particular embodiments, the enhancing agent is acetosyringone, syringaldehyde, methylsyringate or syringic acid.

Other particularly useful enhancing agents may be described by the following formula:



in which formula X represents (-O-) or (-S-), and the substituent groups R^1 - R^9 , which may be identical or different, independently represent any of the following radicals: hydrogen, halogen, hydroxy, formyl, carboxy, and esters and salts hereof, carbamoyl, sulfo, and esters and salts hereof, sulfamoyl, nitro, amino, phenyl, C_1 - C_{14} -alkyl, C_1 - C_5 -alkoxy, carbonyl- C_1 - C_5 -alkyl, aryl- C_1 - C_5 -alkyl; which carbamoyl, sulfamoyl, and amino groups may furthermore be unsubstituted or substituted once or twice with a substituent group R^{10} ; and which phenyl may furthermore be unsubstituted or substituted with one or more substituent groups R^{10} ; and which C_1 - C_{14} -alkyl, C_1 - C_5 -alkoxy, carbonyl- C_1 - C_5 -alkyl, and aryl- C_1 - C_5 -alkyl groups may be saturated or unsaturated, branched or unbranched, and may furthermore be unsubstituted or substituted with one or more substituent groups R^{10} ;

which substituent group R^{10} represents any of the following radicals: halogen, hydroxy, formyl, carboxy and esters and salts hereof, carbamoyl, sulfo and esters and salts hereof, sulfamoyl, nitro, amino, phenyl, aminoalkyl, piperidino, piperazinyl, pyrrolidin-1-yl, C_1 - C_5 -alkyl, C_1 - C_5 -alkoxy; which carbamoyl, sulfamoyl, and amino groups may furthermore be unsubstituted or substituted once or twice with hydroxy, C_1 - C_5 -alkyl, C_1 - C_5 -alkoxy; and which phenyl may furthermore be substituted with one or more of the following radicals: halogen, hydroxy, amino, formyl, carboxy and esters and salts hereof, carbamoyl, sulfo and esters and salts hereof, and sulfamoyl; and which C_1 - C_5 -alkyl, and C_1 - C_5 -alkoxy groups may furthermore be saturated or unsaturated, branched or unbranched, and may furthermore be substituted once or twice with any of the following radicals: halogen, hydroxy, amino, formyl,

carboxy and esters and salts hereof, carbamoyl, sulfo and esters and salts hereof, and sulfamoyl;

or in which general formula two of the substituent groups R^1 - R^9 may together form a group -B-, in which B represents any of the following groups: $(-CHR^{10}-N=N-)$, $(-CH=CH-)_n$, $(-CH=N-)_n$ or $(-N=CR^{10}-NR^{11}-)$, in which groups n represents an integer of from 1 to 3, R^{10} is a substituent group as defined above and R^{11} is defined in the same way as R^{10} . (It is to be understood that if the above mentioned formula comprises two or 10 more R^{10} -substituent groups, these R^{10} -substituent groups may be the same or different).

In particular embodiments, the enhancing agent is 10-methylphenothiazine, phenothiazine-10-propionic acid, N-hydroxysuccinimide phenothiazine-10-propionate, 10-ethylphenothiazine-4-carboxylic acid, 10-ethylphenothiazine, 10-propylphenothiazine, 10-isopropylphenothiazine, methyl phenothiazine-10-propionate, 10-phenylphenothiazine, 10-allylphenothiazine, 10-(3-(4-methylpiperazin-1-yl)propyl)phenothiazine, 10-(2-pyrrolidin-1-yl-ethyl)phenothiazine, 2-methoxy-10-methylphenothiazine, 1-methoxy-10-methylphenothiazine, 3-methoxy-10-methylphenothiazine, 3,10-dimethylphenothiazine, 3,7,10-trimethylphenothiazine, 10-(2-hydroxyethyl)phenothiazine, 10-(3-hydroxypropyl)phenothiazine, 3-(2-hydroxyethyl)-10-methylphenothiazine, 3-hydroxymethyl-10-methylphenothiazine, 25 3,7-dibromophenothiazine-10-propionic acid, phenothiazine-10-propionamide, chlorpromazine, 2-chloro-10-methylphenothiazine, 2-acetyl-10-methylphenothiazine, 10-methylphenoxazine, 10-ethylphenoxazine, phenoxazine-10-propionic acid, 10-(2-hydroxyethyl)phenoxazine or 4-carboxyphenoxazine-10-propionic 30 acid.

The enhancing agents may be obtained from Sigma-Aldrich, Janssen Chimica, Kodak, Tokyo Kasai Organic Chemicals, Daiichi Pure Chemicals Co. or Boehringer Mannheim; N-methylated derivatives of phenothiazine and phenoxazine may be prepared by 35 methylation with methyl iodide as described by Cornel Bodea and Ioan Silberg in "Recent Advances in the Chemistry of Phenothiazines" (Advances in heterocyclic chemistry, 1968, Vol. 9, pp. 321-460); B. Cardillo & G. Casnati in Tetrahedron, 1967,

Vol. 23, p. 3771. Phenothiazine and phenoxazine propionic acids may be prepared as described in J. Org. Chem. 15, 1950, pp. 1125-1130. Hydroxyethyl and hydroxypropyl derivatives of phenothiazine and phenoxazine may be prepared as described by G. Cauquil in Bulletin de la Society Chemique de France, 1960, p.1049.

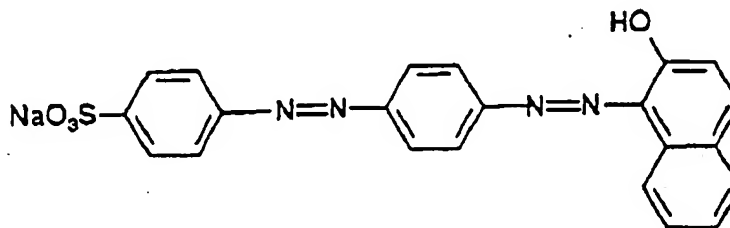
The enhancing agent may be present in concentrations of from 0.01 to 500 μM , preferably in concentrations of from 0.1 to 250 μM .

10 Industrial Applications

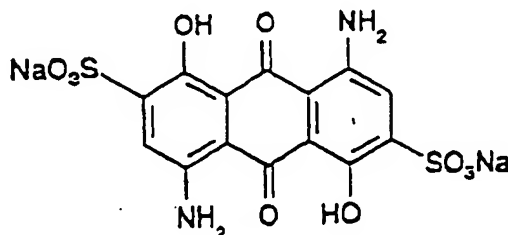
In a preferred embodiment, the method of the invention finds application for bleaching of a dye or dyes in solutions.

The dye may be a synthetic dye such as an azo or an anthraquinone dye, or a natural or nature-identical dye. Examples of dyes are Acid Red 151, Acid blue 45, Direct Blue 1, and indigo carmine, where Acid Red 151, Acid blue 45 and Direct Blue 1 have the following formulas:

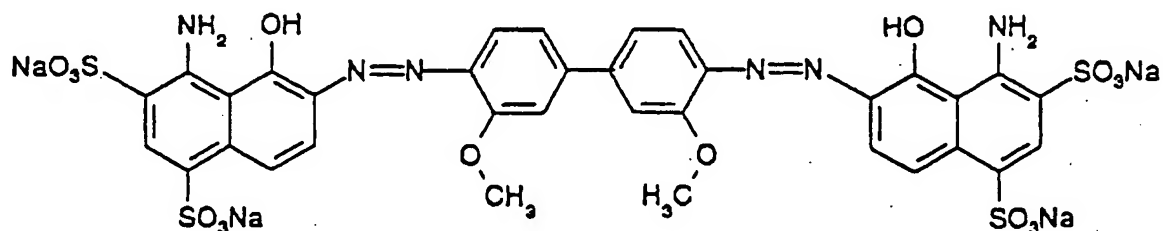
Acid red 151:



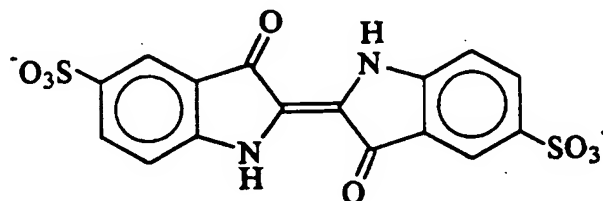
Acid blue 45:



Direct blue 1:



and indigo carmine has the following formula:



In another embodiment, the method of the invention finds application for dye transfer inhibition, e.g., during processing of dyed textiles (cf. e.g. WO 92/18687) or during
5 laundering (cf. e.g. WO 91/05839).

Accordingly, in a specific embodiment, the invention provides a method for inhibiting the transfer of a textile dye from a dyed fabric to another fabric when said fabrics are rinsed together or washed together in a wash liquor, the method
10 comprising treatment of the wash liquor with a laccase obtainable from Coprinaceae in the presence or absence of an enhancing agent.

In another embodiment, the method of the invention finds application in treatment of waste water, e.g., waste
15 water from the chemical or pharmaceutical industry, from dye manufacturing, from dye-works or from the textile industry.

According to the invention the laccase may be present in concentrations of from 0.001-500 LACU/liter, preferably in concentrations of from 0.5-100 LACU/liter.

Detergent Compositions

According to the invention, a laccase of the invention may be added as a component of a detergent composition. As such, it may be included in the detergent composition in the form of a detergent additive. The detergent composition as well as the detergent additive may additionally comprise one or more other enzymes, such as proteases, lipases, amylases, cutinases, cellulases and other oxidoreductases than laccases, e.g., peroxidases and hydrogen peroxide generating oxidases.

10 In a specific aspect, the invention provides a detergent additive. The enzymes may be included in a detergent composition by adding separate additives containing one or more enzymes, or by adding a combined additive comprising all of these enzymes. A detergent additive of the invention, i.e. a separated additive or a combined additive, can be formulated e.g. as granulates, liquids, slurries, etc. Preferred detergent additive formulations are granulates, in particular non-dusting granulates, liquids, in particular stabilized liquids, slurries, or protected enzymes.

20 Non-dusting granulates may be produced, e.g., as disclosed in US 4,106,991 and 4,661,452 (both to Novo Industri A/S) and may optionally be coated by methods known in the art. Examples of waxy coating materials are poly(ethylene oxide) products (polyethyleneglycol, PEG) with mean molecular weights of 1000 to 20000; ethoxylated nonylphenols having from 16 to 50 ethylene oxide units; ethoxylated fatty alcohols in which the alcohol contains from 12 to 20 carbon atoms and in which there are 15 to 80 ethylene oxide units; fatty alcohols; fatty acids; and mono- and di- and triglycerides of fatty acids. Examples of film-forming coating materials suitable for application by fluid bed techniques are given in patent GB 1483591. Liquid enzyme preparations may, for instance, be stabilized by adding a polyol such as propylene glycol, a sugar or sugar alcohol, lactic acid or boric acid according to established methods.

35 Other enzyme stabilizers are well known in the art. Protected enzymes may be prepared according to the method disclosed in EP 238,216.

The detergent composition of the invention may be in any convenient form, e.g., as powder, granules, paste or liquid. A liquid detergent may be aqueous, typically containing up to 70% water and 0-30% organic solvent, or nonaqueous.

5 The detergent composition comprises one or more surfactants, each of which may be anionic, nonionic, cationic, or zwitterionic. The detergent will usually contain 0-50% of anionic surfactant such as linear alkylbenzenesulfonate (LAS), alpha-olefinsulfonate (AOS), alkyl sulfate (fatty alcohol
10 sulfate) (AS), alcohol ethoxysulfate (AEOS or AES), secondary alkanesulfonates (SAS), alpha-sulfo fatty acid methyl esters, alkyl- or alkenylsuccinic acid, or soap. It may also contain 0-40% of nonionic surfactant such as alcohol ethoxylate (AEO or AE), carboxylated alcohol ethoxylates, nonylphenol ethoxylate,
15 alkylpolyglycoside, alkyl dimethylamine oxide, ethoxylated fatty acid monoethanolamide, fatty acid monoethanolamide, or polyhydroxy alkyl fatty acid amide (e.g. as described in WO 92/06154).

The detergent composition may additionally comprise
20 one or more other enzymes, such as amylases, lipases, cutinases, proteases, cellulases, and other oxidoreductases than laccases, e.g., peroxidases and hydrogen peroxide generating oxidases.

The detergent may contain 1-65% of a detergent
25 builder or complexing agent such as zeolite, diphosphate, triphosphate, phosphonate, citrate, nitrilotriacetic acid (NTA), ethylenediaminetetraacetic acid (EDTA), diethylenetriaminepentaacetic acid (DTMPA), alkyl- or alkenylsuccinic acid, soluble silicates or layered silicates (e.g. SKS-6 from
30 Hoechst). The detergent may also be unbuilt, i.e. essentially free of detergent builder.

The detergent may comprise one or more polymers. Examples are carboxymethylcellulose (CMC), poly(vinylpyrrolidone) (PVP), polyethyleneglycol (PEG), poly(vinyl
35 alcohol) (PVA), polycarboxylates such as polyacrylates, maleic/acrylic acid copolymers and lauryl methacrylate/acrylic acid copolymers.

The detergent may contain a bleaching system which may comprise a H_2O_2 source such as perborate or percarbonate which may be combined with a peracid-forming bleach activator such as tetraacetythylenediamine (TAED) or nonanoyloxybenzenesulfonate (NOBS). Alternatively, the bleaching system may comprise peroxyacids of, e.g., the amide, imide, or sulfone type.

The enzymes of the detergent composition of the invention may be stabilized using conventional stabilizing agents, e.g. a polyol such as propylene glycol or glycerol, a sugar or sugar alcohol, lactic acid, boric acid, or a boric acid derivative such as, e.g., an aromatic borate ester, and the composition may be formulated as described in, e.g., WO 92/19709 and WO 92/19708.

The detergent may also contain other conventional detergent ingredients such as, e.g., fabric conditioners including clays, foam boosters, suds suppressors, anti-corrosion agents, soil-suspending agents, anti-soil-redeposition agents, dyes, bactericides, optical brighteners, or perfume.

The pH (measured in aqueous solution at use concentration) will usually be neutral or alkaline, e.g. in the range of 7-11.

Particular forms of detergent compositions within the scope of the invention include:

1) A detergent composition formulated as a granulate having a bulk density of at least 600 g/l comprising

	Linear alkylbenzenesulfonate (calculated as acid)	7	-	12%
30	Alcohol ethoxysulfate (e.g. C_{12-18} alcohol, 1-2 EO) or alkyl sulfate (e.g. C_{16-18})	1	-	4%
	Alcohol ethoxylate (e.g. C_{14-15} alcohol, 7 EO)	5	-	9%
35	Sodium carbonate (as Na_2CO_3)	14	-	20%
	Soluble silicate (as $Na_2O, 2SiO_2$)	2	-	6%
	Zeolite (as $NaAlSiO_4$)	15	-	22%

	Sodium sulfate (as Na_2SO_4)	0 - 6%
	Sodium citrate/citric acid (as $\text{C}_6\text{H}_5\text{Na}_3\text{O}_7/\text{C}_6\text{H}_8\text{O}_7$)	0 - 15%
	Sodium perborate (as $\text{NaBO}_3 \cdot \text{H}_2\text{O}$)	11 - 18%
5	TAED	2 - 6%
	Carboxymethylcellulose	0 - 2%
	Polymers (e.g. maleic/acrylic acid copolymer, PVP, PEG)	0 - 3%
10	Enzymes (calculated as pure enzyme protein)	0.0001 - 0.1%
	Minor ingredients (e.g. suds suppressors, perfume, optical brightener, photobleach)	0 - 5%

2) A detergent composition formulated as a granulate having a
15 bulk density of at least 600 g/l comprising

	Linear alkylbenzenesulfonate (cal- culated as acid)	6 - 11%
20	Alcohol ethoxysulfate (e.g. C_{12-18} alcohol, 1-2 EO or alkyl sulfate (e.g. C_{16-18}))	1 - 3%
	Alcohol ethoxylate (e.g. C_{14-15} alco- hol, 7 EO)	5 - 9%
	Sodium carbonate (as Na_2CO_3)	15 - 21%
25	Soluble silicate (as $\text{Na}_2\text{O} \cdot 2\text{SiO}_2$)	1 - 4%
	Zeolite (as NaAlSiO_4)	24 - 34%
	Sodium sulfate (as Na_2SO_4)	4 - 10%
	Sodium citrate/citric acid (as $\text{C}_6\text{H}_5\text{Na}_3\text{O}_7/\text{C}_6\text{H}_8\text{O}_7$)	0 - 15%
30	Carboxymethylcellulose	0 - 2%
	Polymers (e.g. maleic/acrylic acid copolymer, PVP, PEG)	1 - 6%
	Enzymes (calculated as pure enzyme protein)	0.0001 - 0.1%
35	Minor ingredients (e.g. suds suppressors, perfume)	0 - 5%

3) A detergent composition formulated as a granulate having a bulk density of at least 600 g/l comprising

	Linear alkylbenzenesulfonate (calculated as acid)	5	-	9%
5	Alcohol ethoxylate (e.g. C ₁₂₋₁₅ alcohol, 7 EO)	7	-	14%
	Soap as fatty acid (e.g. C ₁₆₋₂₂ fatty acid)	1	-	3%
10	Sodium carbonate (as Na ₂ CO ₃)	10	-	17%
	Soluble silicate (as Na ₂ O, 2SiO ₂)	3	-	9%
	Zeolite (as NaAlSiO ₄)	23	-	33%
	Sodium sulfate (as Na ₂ SO ₄)	0	-	4%
	Sodium perborate (as NaBO ₃ ·H ₂ O)	8	-	16%
15	TAED	2	-	8%
	Phosphonate (e.g. EDTMPA)	0	-	1%
	Carboxymethylcellulose	0	-	2%
	Polymers (e.g. maleic/acrylic acid copolymer, PVP, PEG)	0	-	3%
20	Enzymes (calculated as pure enzyme protein)	0.0001	-	0.1%
	Minor ingredients (e.g. suds suppressors, perfume, optical brightener)	0	-	5%

25 4) A detergent composition formulated as a granulate having a bulk density of at least 600 g/l comprising

	Linear alkylbenzenesulfonate (calculated as acid)	8	-	12%
30	Alcohol ethoxylate (e.g. C ₁₂₋₁₅ alcohol, 7 EO)	10	-	25%
	Sodium carbonate (as Na ₂ CO ₃)	14	-	22%
	Soluble silicate (as Na ₂ O, 2SiO ₂)	1	-	5%
	Zeolite (as NaAlSiO ₄)	25	-	35%
35	Sodium sulfate (as Na ₂ SO ₄)	0	-	10%
	Carboxymethylcellulose	0	-	2%

	Polymers (e.g. maleic/acrylic acid copolymer, PVP, PEG)	1	-	3%
	Enzymes (calculated as pure enzyme protein)	0.0001	-	0.1%
5	Minor ingredients (e.g. suds suppressors, perfume)	0	-	5%

5) An aqueous liquid detergent composition comprising

	Linear alkylbenzenesulfonate (calculated as acid)	15	-	21%
10	Alcohol ethoxylate (e.g. C ₁₂₋₁₅ alcohol, 7 EO or C ₁₂₋₁₅ alcohol, 5 EO)	12	-	18%
	Soap as fatty acid (e.g. oleic acid)	3	-	13%
15	Alkenylsuccinic acid (C ₁₂₋₁₄)	0	-	13%
	Aminoethanol	8	-	18%
	Citric acid	2	-	8%
	Phosphonate	0	-	3%
	Polymers (e.g. PVP, PEG)	0	-	3%
20	Borate (as B ₄ O ₇)	0	-	2%
	Ethanol	0	-	3%
	Propylene glycol	8	-	14%
	Enzymes (calculated as pure enzyme protein)	0.0001	-	0.1%
25	Minor ingredients (e.g. dispersants, suds suppressors, perfume, optical brightener)	0	-	5%

6) An aqueous structured liquid detergent composition comprising

	Linear alkylbenzenesulfonate (calculated as acid)	15	- 21%
5	Alcohol ethoxylate (e.g. C ₁₂₋₁₅ alcohol, 7 EO, or C ₁₂₋₁₅ alcohol, 5 EO)	3	- 9%
	Soap as fatty acid (e.g. oleic acid)	3	- 10%
10	Zeolite (as NaAlSiO ₄)	14	- 22%
	Potassium citrate	9	- 18%
	Borate (as B ₄ O ₇)	0	- 2%
	Carboxymethylcellulose	0	- 2%
	Polymers (e.g. PEG, PVP)	0	- 3%
15	Anchoring polymers such as, e.g., lauryl methacrylate/acrylic acid copolymer; molar ratio 25:1; MW 3800	0	- 3%
	Glycerol	0	- 5%
20	Enzymes (calculated as pure enzyme protein)	0.0001	- 0.1%
	Minor ingredients (e.g. dispersants, suds suppressors, perfume, optical brighteners)	0	- 5%

25 7) A detergent composition formulated as a granulate having a
bulk density of at least 600 g/l comprising

	Fatty alcohol sulfate	5	- 10%
	Ethoxylated fatty acid monoethanol- amide	3	- 9%
30	Soap as fatty acid	0	- 3%
	Sodium carbonate (as Na ₂ CO ₃)	5	- 10%
	Soluble silicate (as Na ₂ O, 2SiO ₂)	1	- 4%
	Zeolite (as NaAlSiO ₄)	20	- 40%
	Sodium sulfate (as Na ₂ SO ₄)	2	- 8%
35	Sodium perborate (as NaBO ₃ ·H ₂ O)	12	- 18%
	TAED	2	- 7%

	Polymers (e.g. maleic/acrylic acid copolymer, PEG)	1	-	5%
	Enzymes (calculated as pure enzyme protein)	0.0001	-	0.1%
5	Minor ingredients (e.g. optical brightener, suds suppressors, perfume)	0	-	5%

8) A detergent composition formulated as a granulate comprising

10	Linear alkylbenzenesulfonate (calculated as acid)	8	-	14%
	Ethoxylated fatty acid monoethanolamide	5	-	11%
	Soap as fatty acid	0	-	3%
	Sodium carbonate (as Na_2CO_3)	4	-	10%
15	Soluble silicate (as $\text{Na}_2\text{O}, 2\text{SiO}_2$)	1	-	4%
	Zeolite (as NaAlSiO_4)	30	-	50%
	Sodium sulfate (as Na_2SO_4)	3	-	11%
	Sodium citrate (as $\text{C}_6\text{H}_5\text{Na}_3\text{O}_7$)	5	-	12%
20	Polymers (e.g. PVP, maleic/acrylic acid copolymer, PEG)	1	-	5%
	Enzymes (calculated as pure enzyme protein)	0.0001	-	0.1%
	Minor ingredients (e.g. suds suppressors, perfume)	0	-	5%

25 9) A detergent composition formulated as a granulate comprising

	Linear alkylbenzenesulfonate (calculated as acid)	6	-	12%
	Nonionic surfactant	1	-	4%
	Soap as fatty acid	2	-	6%
30	Sodium carbonate (as Na_2CO_3)	14	-	22%
	Zeolite (as NaAlSiO_4)	18	-	32%
	Sodium sulfate (as Na_2SO_4)	5	-	20%
	Sodium citrate (as $\text{C}_6\text{H}_5\text{Na}_3\text{O}_7$)	3	-	8%
	Sodium perborate (as $\text{NaBO}_3 \cdot \text{H}_2\text{O}$)	4	-	9%
35	Bleach activator (e.g. NOBS or TAED)	1	-	5%

	Carboxymethylcellulose	0	-	2%
	Polymers (e.g. polycarboxylate or PEG)	1	-	5%
5	Enzymes (calculated as pure enzyme protein)	0.0001	-	0.1%
	Minor ingredients (e.g. optical brightener, perfume)	0	-	5%

10) An aqueous liquid detergent composition comprising

10	Linear alkylbenzenesulfonate (calculated as acid)	15	-	23%
	Alcohol ethoxysulfate (e.g. C ₁₂₋₁₅ alcohol, 2-3 EO)	8	-	15%
15	Alcohol ethoxylate (e.g. C ₁₂₋₁₅ alcohol, 7 EO, or C ₁₂₋₁₅ alcohol, 5 EO)	3	-	9%
	Soap as fatty acid (e.g. lauric acid)	0	-	3%
	Aminoethanol	1	-	5%
	Sodium citrate	5	-	10%
20	Hydrotrope (e.g. sodium toluenesulfonate)	2	-	6%
	Borate (as B ₄ O ₇)	0	-	2%
	Carboxymethylcellulose	0	-	1%
	Ethanol	1	-	3%
25	Propylene glycol	2	-	5%
	Enzymes (calculated as pure enzyme protein)	0.0001	-	0.1%
30	Minor ingredients (e.g. polymers, dispersants, perfume, optical brighteners)	0	-	5%

11) An aqueous liquid detergent composition comprising

	Linear alkylbenzenesulfonate (calculated as acid)	20	-	32%
35	Alcohol ethoxylate (e.g. C ₁₂₋₁₅ alcohol, 7 EO, or C ₁₂₋₁₅ alcohol, 5 EO)	6	-	12%
	Aminoethanol	2	-	6%

	Citric acid	8	- 14%
	Borate (as B_4O_7)	1	- 3%
5	Polymer (e.g. maleic/acrylic acid copolymer, anchoring polymer such as, e.g., lauryl methacrylate/acrylic acid copolymer)	0	- 3%
	Glycerol	3	- 8%
10	Enzymes (calculated as pure enzyme protein)	0.0001	- 0.1%
	Minor ingredients (e.g. hydro-tropes, dispersants, perfume, optical brighteners)	0	- 5%

12) A detergent composition formulated as a granulate having
 15 a bulk density of at least 600 g/l comprising

20	Anionic surfactant (linear alkylbenzenesulfonate, alkyl sulfate, alpha-olefinsulfonate, alpha-sulfo fatty acid methyl esters, alkanesulfonates, soap)	25	- 40%
	Nonionic surfactant (e.g. alcohol ethoxylate)	1	- 10%
	Sodium carbonate (as Na_2CO_3)	8	- 25%
	Soluble silicates (as Na_2O , $2SiO_2$)	5	- 15%
25	Sodium sulfate (as Na_2SO_4)	0	- 5%
	Zeolite (as $NaAlSiO_4$)	15	- 28%
	Sodium perborate (as $NaBO_3 \cdot 4H_2O$)	0	- 20%
	Bleach activator (TAED or NOBS)	0	- 5%
30	Enzymes (calculated as pure enzyme protein)	0.0001	- 0.1%
	Minor ingredients (e.g. perfume, optical brighteners)	0	- 3%

13) Detergent formulations as described in 1) - 12) wherein all
 or part of the linear alkylbenzenesulfonate is replaced by (C_{12} -
 35 C_{18}) alkyl sulfate.

14) A detergent composition formulated as a granulate having
 a bulk density of at least 600 g/l comprising

	(C ₁₂ -C ₁₈) alkyl sulfate	9	- 15%
	Alcohol ethoxylate	3	- 6%
	Polyhydroxy alkyl fatty acid amide	1	- 5%
	Zeolite (as NaAlSiO ₄)	10	- 20%
5	Layered disilicate (e.g. SK56 from Hoechst)	10	- 20%
	Sodium carbonate (as Na ₂ CO ₃)	3	- 12%
	Soluble silicate (as Na ₂ O, 2SiO ₂)	0	- 6%
	Sodium citrate	4	- 8%
10	Sodium percarbonate	13	- 22%
	TAED	3	- 8%
	Polymers (e.g. polycarboxylates and PVP)	0	- 5%
15	Enzymes (calculated as pure enzyme protein)	0.0001	- 0.1%
	Minor ingredients (e.g. optical brightener, photo bleach, perfume, suds suppressors)	0	- 5%

15) A detergent composition formulated as a granulate having
 20 a bulk density of at least 600 g/l comprising

	(C ₁₂ -C ₁₈) alkyl sulfate	4	- 8%
	Alcohol ethoxylate	11	- 15%
	Soap	1	- 4%
	Zeolite MAP or zeolite A	35	- 45%
25	Sodium carbonate (as Na ₂ CO ₃)	2	- 8%
	Soluble silicate (as Na ₂ O, 2SiO ₂)	0	- 4%
	Sodium percarbonate	13	- 22%
	TAED	1	- 8%
	Carboxymethyl cellulose	0	- 3%
30	Polymers (e.g. polycarboxylates and PVP)	0	- 3%
	Enzymes (calculated as pure enzyme protein)	0.0001	- 0.1%
35	Minor ingredients (e.g. optical brightener, phosphonate, perfume)	0	- 3%

- 16) Detergent formulations as described in 1) - 15) which contain a stabilized or encapsulated peracid, either as an additional component or as a substitute for already specified bleach systems.
- 5 17) Detergent compositions as described in 1), 3), 7), 9) and 12) wherein perborate is replaced by percarbonate.
- 18) Detergent compositions as described in 1), 3), 7), 9), 12), 14) and 15) which additionally contain a manganese catalyst. The manganese catalyst may, e.g., be one of the compounds
10 described in "Efficient manganese catalysts for low-temperature bleaching", Nature 369, 1994, pp. 637-639.
- 19) Detergent composition formulated as a nonaqueous detergent liquid comprising a liquid nonionic surfactant such as, e.g., linear alkoxylated primary alcohol, a builder system (e.g.
15 phosphate), enzyme and alkali. The detergent may also comprise anionic surfactant and/or a bleach system.

The laccases of the invention may be incorporated at an enzyme protein level conventionally employed for other enzymes in detergents. It is at present contemplated that, in detergent
20 compositions of the invention, the laccases may be added at a level corresponding to 0.00001-5 mg, preferably at a level corresponding to 0.0001-1 mg (calculated as pure enzyme protein) per liter of wash liquor.

The present invention is further illustrated in the
25 following examples which are not in any way intended to limit the scope of the invention as claimed.

EXAMPLE 1

Bleaching of Direct Blue 1 (DB1) using Coprinaceae laccases with various enhancing agents at pH 5.5-8.5.

Bleaching of the dye Direct Blue 1 at various pH values was conducted using a laccase obtained from Coprinus comatus, Coprinus friesii, Coprinus plicatilis, Panaeolus papilionaceus or Psathyrella condolleana and one of the following enhancing agents:

- PPT (phenothiazine-10-propionic acid)
 10 PPO (phenoxazine-10-propionic acid)
 methylsyringate.

The above mentioned strains were fermented in the following way:

The strains were inoculated on PDA agar plates (PDA: 39 g/1 potato dextrose agar) and grown at 26°C for 3 days. Shake flasks were then inoculated with 6-8 small squares (0.5 cm x 0.5 cm) of agar containing mycelium and fermented for 3-10 days at 26°C and 200 rpm using the following medium:

	Deposit no.	Medium	Growth
20 <u>Coprinus comatus</u>	CBS 631.95	A	10 days
<u>Coprinus friesii</u>	CBS 629.95	A	3 days
<u>Panaeolus papilionaceus</u>	CBS 630.95	A	10 days
<u>Psathyrella condolleana</u>	CBS 628.95	B	7 days
25 <u>Coprinus plicatilis</u>	CBS 627.95	A	8 days

Media:

A:	soja meal	30 g/l
	maltodextrin	15 g/l
30	bacto peptone	5 g/l
	pluronic	0.2 g/l

B:	potato meal	50 g/l
	barley meal	25 g/l
	BAN 800MG*	0.025 g/l
	Na-caseinate	5 g/l
5	crushed soja	10 g/l
	Na ₂ HPO ₄ , 12 H ₂ O	4.5 g/l
	Pluronic	0.05 ml/l
* BAN 800MG obtainable from Novo Nordisk A/S.		

After fermentation the culture broths were centrifugated
10 and the supernatants were used in the tests described below.

The bleaching rate of DB1 was determined using the
following conditions:

	Final concentration
400 µl 50 mM Britton-Robinson buffer*,	
15 (pH 5.5, 7.0, and 8.5 respectively),	20 mM
200 µl DB1 ~ 3.0 Abs. Units (610 nm)	0.6 (A _{610nm})
200 µl 50 µM enhancing agent	10 µM
200 µl laccase	4 LACU/l at pH
	5.5 and at pH 7.0; 20 LACU/l at pH 8.5.

20 * (50 mM acetic acid, 50 mM phosphoric acid, 50 mM boric acid,
pH adjusted to the value of interest with NaOH).

Reagents were mixed in a 1 ml thermostated cuvette at
30°C and the bleaching was started by addition of the laccase.

The bleaching was followed spectrophotometrically at 610
25 nm, which is the wavelength of the absorption peak of DB1, with
readings every 5 sec. for a period of 5 minutes. The initial
bleaching rate was determined from the first linear part of the
absorbance curve.

The following results were obtained with PPT:

		-ΔmAbs/minute		
Laccase:				
	pH:	5.5	7.0	8.5
5				
	C. comatus	68	17	3
	C. friesii	221	44	36
	Pan. papilionaceus	91	25	9
	Ps. condolleana	88	36	25
10	C. pliccalitis	75	33	0

The following results were obtained with PPO:

		-ΔmAbs/minute		
Laccase:				
	pH:	5.5	7.0	8.5
15				
	C. comatus	102	44	4
	C. friesii	122	80	116
	Pan. papilionaceus	120	67	47
	Ps. condolleana	114	114	72
20	C. pliccalitis	106	83	23

The following results were obtained with methyl-syringate:

		-ΔmAbs/minute		
Laccase:				
	pH:	5.5	7.0	8.5
25				
	C. comatus	33	23	2
	C. friesii	40	55	61
	Pan. papilionaceus	16	19	18
30	Ps. condolleana	45	54	43
	C. pliccalitis	42	39	14

EXAMPLE 2Bleaching of Direct Blue 1 (DB1) using *Coprinus cinereus* laccase with/without enhancing agents at pH 5.5-8.5.

Bleaching of the dye Direct Blue 1 at various pH values
5 was conducted using *Coprinus cinereus* laccase and one of the
following enhancing agents:

None

PPT (phenothiazine-10-propionic acid)

MPT (10-methylphenothiazine)

10 PPO (phenoxazine-10-propionic acid)

MPO (10-methylphenoxazine)

EPC (10-ethylphenothiazine-4-carboxylic acid)

acetosyringone

syringaldehyde

15 methylsyringate

ABTS (2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonate)).

The laccase was obtained in the following way:

Coprinus cinereus (IFO 30116) was inoculated from a PDA agar
slant (PDA: 39 g/l potato dextrose agar) into a 100 ml shake
20 flask containing medium A (Medium A is described in Example 1).
The culture was cultivated for 6 days at 26°C and 100 rpm. A
10-liter fermentor containing medium A was inoculated with the
100 ml culture broth. The fermentation ran for 6 days at 26°C
and 100 rpm. The culture broth was filtrated and concentrated
25 by ultrafiltration. Further purification was carried out using
hydrophobic interaction chromatography followed by anionic
exchange chromatography. This process resultated in a
preparation with a laccase activity of 3.6 LACU/ml. The
estimated purity was >80% on a protein basis.

30 The bleaching rate of DB1 was determined using the
following conditions:

	Final concentration
400 μ l 50 mM Britton-Robinson buffer*, (pH 5.5, 7.0 and 8.5 respectively),	20 mM
200 μ l DB1 ~ 3.0 Abs. Units (610 nm)	0.6 (A_{610nm})
5 200 μ l 50 μ M enhancing agent	10 μ M
200 μ l <u>C. cinereus</u> laccase	4 LACU/l

* (50 mM acetic acid, 50 mM phosphoric acid, 50 mM boric acid, pH adjusted to the value of interest with NaOH).

Reagents were mixed in a 1 ml thermostated cuvette at 10 30°C and the bleaching was started by addition of the laccase.

The bleaching was followed spectrophotometrically at 610 nm, which is the wavelength of the absorption peak of DB1, with readings every 5 sec. for a period of 5 minutes. The initial bleaching rate was determined from the first linear part of the 15 absorbance curve.

The following results were obtained:

Enhancing agent	-ΔmAbs/minute		
pH:	5.5	7.0	8.5
20 none	13	5	3
PPT	74	74	24
MPT	43	30	1
PPO	121	104	48
MPO	83	74	30
25 EPC	184	122	57
aceto-			
syringone	28	94	50
syring-			
aldehyde	29	79	28
30 methyl-			
syringate	20	94	57
ABTS	84	76	24

EXAMPLE 3Bleaching of Direct Blue 1 (DB1) using *Coprinus cinereus* laccase and acetosyringone

Bleaching of the dye Direct Blue 1 at various pH values was conducted using *Coprinus cinereus* laccase and the enhancing agent acetosyringone.

The laccase was obtained as described in Example 2. Acetosyringone was obtained from Aldrich.

The bleaching rate of DB1 was determined using the following conditions:

	Final concentration
400 μ l 50 mM Britton-Robinson buffer*, (pH 4, 5, 6, 7, and 8 respectively),	20 mM
200 μ l DB1 ~ 3.0 Abs. Units (610 nm)	0.6 (A_{610nm})
15 200 μ l 50 μ M acetosyringone	10 μ M
200 μ l <u><i>C. cinereus</i></u> laccase	13 LACU/l

* (50 mM acetic acid, 50 mM phosphoric acid, 50 mM boric acid, pH adjusted to the value of interest with NaOH).

Reagents were mixed in a 1 cm thermostated cuvette at 20 30°C and the bleaching was started by addition of the laccase.

The bleaching was detected spectrophotometrically at 610 nm, which is the wavelength of the absorption peak of DB1. After 5 sec. bleaching was followed for 4 minutes.

The following results were obtained:

25 pH	Initial DB1 bleaching ($-\Delta mAbs/\text{min}$) (% of pH 7-value)
-------	---

4	18 %
30 5	13 %
6	35 %
7	100 %
8	69 %

It can be seen from the results given above that the optimum bleaching is achieved at pH around 7, but the system also shows a significant bleaching at pH 8.

EXAMPLE 4

5 Bleaching of gradually added Acid Blue 45 with Coprinus cinereus Laccase with and without enhancing agents

Ideally, dye transfer inhibition systems for laundry applications should be tested in a real wash where dyed fabrics give off dyes to the wash solution as a result of the combined
10 action of the detergent, temperature and mechanical agitation taking place.

To simulate such a process, however, a magnetically stirred beaker was used as the reaction vessel and dye was added gradually from a stock solution (using a Metrohm 725
15 dosimat). The solution was monitored spectrophotometrically using a Zeiss multichannel spectrometer (MCS) equipped with a fibre-optics immersion probe.

Stock solutions of the enhancing agents were prepared either in a suitable water/ethanol mixture or by first dissol-
20 ving the enhancing agent in a small amount of dimethylformamide and then diluting with a pH 10.0 phosphate/borate buffer. Stock solutions of the dye Acid Blue 45 were made with water.

The laccase was obtained as described in Example 2.

The following conditions were used in the experiment:

- 25 Temperature: 35°C
- Medium and pH: 50 mM/50 mM phosphate/borate buffer at pH 10
- Enhancing agent (when applicable): 10 μ M
- Laccase: 0.04 LACU/ml
- 30 Dye addition program: linear addition at rates of ca 0.34 abs/40 min, referring to the absorbance of the dye at its maximum absorbance wavelength (590 nm for Acid Blue 45).

The following enhancing agents were tested: phenoxazine-10-propionic acid and 10-ethylphenothiazine-4-carboxylic acid.

Fig. 1 shows the results of the bleaching tests. The following symbols are used: (I): Only dye addition; (II): Dye addition in the presence of Laccase; (III): Dye addition in the presence of Laccase + 10-ethylphenothiazine-4-carboxylic acid; 5 (IV): Dye addition in the presence of Laccase + phenoxazine-10-propionic acid.

It can be seen from Fig. 1 that the tested laccase performs well even without enhancing agents at pH 10; but the bleaching effect is enhanced by phenoxazine-10-propionic acid 10 and 10-ethylphenothiazine-4-carboxylic acid.

EXAMPLE 5

Dye Transfer Inhibition Using Coprinus cinereus Laccase

A small-scale experiment was carried out in which clean cotton test pieces were washed together with dyed fabrics 15 bleeding dye into the wash solution, the experiment conducted in the absence and in the presence of laccase and enhancing agent.

After wash, the Hunter colour difference between the above mentioned cotton pieces and clean cotton pieces (washed 20 in the absence of bleeding fabrics) was measured and taken as a measure of the degree of dye transfer resulting from the wash.

Materials used:

Bleeding fabrics dyed with Acid Red 151 (AR 151) or 25 Direct Blue 1 (DB1).

Clean white cotton (bleached, no optical brightener added).

Liquid detergent and powder detergent as typically met in the North American market place; both detergents contained 30 no bleaching system.

Coprinus cinereus laccase, obtained as described in Example 2.

Washing procedure:

The washing processes were carried out in beakers with magnetical stirring at 35°C for 15 min., after which the test fabrics were rinsed thoroughly in tap water and air-dried overnight in the dark before the Hunter readings were taken by using a Datacolor Elrephometer 2000 reflectance spectrometer.

Treatment 1 was in each case a wash with laccase at a level of 40 LACU/l with the enhancing agent 10-ethylphenothiazine-4-carboxylic acid at a level of 10 µM.

10 Treatment 2 was in each case a wash with laccase at a level of 40 LACU/l with the enhancing agent acetosyringone at a level of 10 µM.

The following results were obtained:

15 Wash in liquid detergent solution (2 g/l, water hardness 6°dH) at pH 8.5:

Hunter colour difference (delta E)
with respect to white, washed cotton

	Cotton washed with AR 151	Cotton washed with DB 1
20 Wash with no laccase system	12	26
Treatment 1	9	3
Treatment 2	1	7

25 Wash in powder detergent solution (1 g/l, water hardness 6°dH) at pH 10.0:

Hunter colour difference (delta E)
with respect to white, washed cotton

Cotton washed with AR 151	Cotton washed with DB 1
------------------------------	----------------------------


Wash with no laccase		
system	21	29
Treatment 1	17	13
5 Treatment 2	4	8

Typical significant differences in the delta E readings are 2-3 units, so the data reflect significant reduction of dye transfer with the laccase treatments relative to the treatment with no laccase system.

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>3</u> , lines <u>11-16</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution CENTRAALBUREAU VOOR SCHIMMELCULTURES	
Address of depositary institution (including postal code and country) Oosterstraat 1, Postbus 273, NL-3740 AG Barn, Nether- land	
Date of deposit 16 August 1995	Accession Number CBS 631.95
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect of those designations in which a European and/or Australian patent is sought, during the pendency of the patent application a sample of the deposited microorganism is only to be provided to an independent expert nominated by the person requesting the sample (Rule 28(4) EPC / Regulation 3.25 of Australia Statutory Rules 1991 No 71).	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="checked" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer 	Authorized officer

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>3</u> , line S <u>17-22</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution CENTRAALBUREAU VOOR SCHIMMELCULTURES	
Address of depositary institution (including postal code and country) Oosterstraat 1, Postbus 273, NL-3740 AG Barn, Nether- land	
Date of deposit 16 August 1995	Accession Number CBS 629.95
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect of those designations in which a European and/or Australian patent is sought, during the pendency of the patent application a sample of the deposited microorganism is only to be provided to an independent expert nominated by the person requesting the sample (Rule 28(4) EPC / Regulation 3.25 of Australia Statutory Rules 1991 No 71).	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	

For receiving Office use only

☒ This sheet was received with the international application

Authorized officer



For International Bureau use only

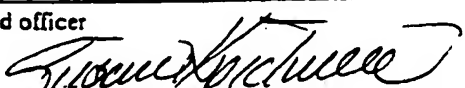
☐ This sheet was received by the International Bureau on:

Authorized officer

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>3</u> , lineS <u>23-29</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution CENTRAALBUREAU VOOR SCHIMMELCULTURES	
Address of depositary institution (including postal code and country) Oosterstraat 1, Postbus 273, NL-3740 AG Barn, Nether- land	
Date of deposit 16 August 1995	Accession Number CBS 627.95
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect of those designations in which a European and/or Australian patent is sought, during the pendency of the patent application a sample of the deposited microorganism is only to be provided to an independent expert nominated by the person requesting the sample (Rule 28(4) EPC / Regulation 3.25 of Australia Statutory Rules 1991 No 71).	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	


For receiving Office use only
<input checked="" type="checkbox"/> This sheet was received with the international application
Authorized officer 

For International Bureau use only
<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>3</u> , line <u>30-36</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution <p style="text-align: center;">CENTRAALBUREAU VOOR SCHIMMELCULTURES</p>	
Address of depositary institution (including postal code and country) <p style="text-align: center;">Oosterstraat 1, Postbus 273, NL-3740 AG Barn, Nether- land</p>	
Date of deposit <p style="text-align: center;">16 August 1995</p>	Accession Number <p style="text-align: center;">CBS 628.95</p>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
<p>In respect of those designations in which a European and/or Australian patent is sought, during the pendency of the patent application a sample of the deposited microorganism is only to be provided to an independent expert nominated by the person requesting the sample (Rule 28(4) EPC / Regulation 3.25 of Australia Statutory Rules 1991 No 71).</p>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	


For receiving Office use only
<input checked="" type="checkbox"/> This sheet was received with the international application
Authorized officer 

For International Bureau use only
<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page S <u>3 and 4</u> , line <u>37-38 (page 3) and 1-5 (page 4)</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution CENTRAALBUREAU VOOR SCHIMMELCULTURES	
Address of depositary institution (including postal code and country) Oosterstraat 1, Postbus 273, NL-3740 AG Barn, Nether- land	
Date of deposit 16 August 1995	Accession Number CBS 630.95
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
In respect of those designations in which a European and/or Australian patent is sought, during the pendency of the patent application a sample of the deposited microorganism is only to be provided to an independent expert nominated by the person requesting the sample (Rule 28(4) EPC / Regulation 3.25 of Australia Statutory Rules 1991 No 71).	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	

For receiving Office use only
<input checked="" type="checkbox"/> This sheet was received with the international application
Authorized officer 

For International Bureau use only
<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer

CLAIMS

1. A method of oxidizing a substrate in a solution with a pH at or above 7, comprising contacting the substrate with an effective amount of a laccase obtainable from Coprinaceae.

2. A method according to claim 1, wherein the laccase is obtainable from Coprinus, Podaxis, Montagnea, Macrometrula, Psathyrella, Panaeolina, Panaeolus, Copelandia, Anellaria, Limnoperdon, Panaeolopsis or Polyplocium.

3. A method according to claim 2, wherein the laccase is obtainable from Coprinus cinereus.

4. A method according to claim 3, wherein the laccase is obtainable from Coprinus cinereus IFO30116, or is a laccase displaying immunochemical cross-reactivity with an antibody raised against a laccase obtainable from Coprinus cinereus IFO30116.

5. A method according to claim 1, wherein an effective amount of laccase is in the range of from 0.001 LACU/liter to 500 LACU/liter.

6. A method according to claims 1-5, additionally comprising contacting the substrate with an effective amount of an enhancing agent.

7. A method according to claim 6, wherein the enhancing agent is phenothiazine or phenoxazine or a derivative one of these.

8. A method according to claim 7, wherein the enhancing agent is selected from the group consisting of phenothiazine-10-propionic acid, 10-methylphenothiazine,

phenoxazine-10-propionic acid, 10-methylphenoxazine and 10-ethylphenothiazine-4-carboxylic acid.

9. A method according to claim 6, wherein the enhancing agent is acetosyringone, syringaldehyde or methyl-
5 syringate.

10. A method according to any of claims 1-9, in which said method is a method for bleaching of a dye or dyes in solution.

11. A method according to any of claims 1-10, in
10 which said method is a method for inhibiting the transfer of a textile dye from a dyed fabric to another fabric when said fabrics are washed together in a wash liquor.

12. A method according to any of claims 6-9, in which the enhancing agent is added at the beginning of or during the
15 process.

13. A method according to any preceding claim, in which the level of enhancing agent added is in the range of from 0.01-500 μ M, preferably in the range of from 0.1-250 μ M.

14. A detergent additive for inhibiting the transfer
20 of a textile dye from a dyed fabric to another fabric when said fabrics are washed and/or rinsed together in a wash liquor characterized by comprising

- a) a laccase obtainable from Coprinaceae
- b) optionally an enhancing agent.

25 15. A detergent additive according to claim 14, wherein the laccase is obtainable from Coprinus, in particular from Coprinus cinereus.

16. A detergent additive according to claims 14-15, wherein the laccase is obtainable from Coprinus cinereus
30 IF030116, or is a laccase displaying immunochemical cross-

reactivity with an antibody raised against a laccase obtainable from Coprinus cinereus IF030116.

17. A detergent additive according to claims 14-16, wherein the enhancing agent is phenothiazine or phenoxazine or a derivative one of these.

18. A detergent additive according to claim 17, wherein the enhancing agent is selected from the group consisting of phenothiazine-10-propionic acid, 10-methylphenothiazine, phenoxazine-10-propionic acid, 10-methylphenoxazine and 10-ethylphenothiazine-4-carboxylic acid.

19. A detergent additive according to claims 14-16, wherein the enhancing agent is acetosyringone, syringaldehyde or methylsyringate.

20. A detergent additive according to any of claims 14-19, provided in the form of a granulate, preferably a non-dusting granulate, a liquid, in particular a stabilized liquid, a slurry, or a protected enzyme.

21. A detergent composition comprising a laccase obtainable from Coprinaceae, a surfactant and optionally an enhancing agent.

22. A detergent composition according to claim 21, wherein the laccase is obtainable from Coprinus, in particular from Coprinus cinereus.

23. A detergent composition according to claims 21-22, wherein the laccase is obtainable from Coprinus cinereus IF030116 or is a laccase displaying immunochemical cross-reactivity with an antibody raised against a laccase derived from Coprinus cinereus IF030116.

24. A detergent composition according to claims 21-23, wherein the enhancing agent is phenothiazine or phenoxazine or a derivative one of these.

25. A detergent composition according to claim 24,
5 wherein the enhancing agent is selected from the group consisting of phenothiazine-10-propionic acid, 10-methylphenothiazine, phenoxazine-10-propionic acid, 10-methylphenoxazine and 10-ethylphenothiazine-4-carboxylic acid.

26. A detergent composition according to claim 23,
10 wherein the enhancing agent is acetosyringone, syringaldehyde or methylsyringate.

27. A detergent composition according to any of claims 21-26, which further comprises one or more other enzymes, in particular a protease, a lipase, an amylase, a
15 cutinase, a cellulase and/or an another oxidoreductase than laccase, e.g., a peroxidase and/or a hydrogen peroxide generating oxidase.

1/1

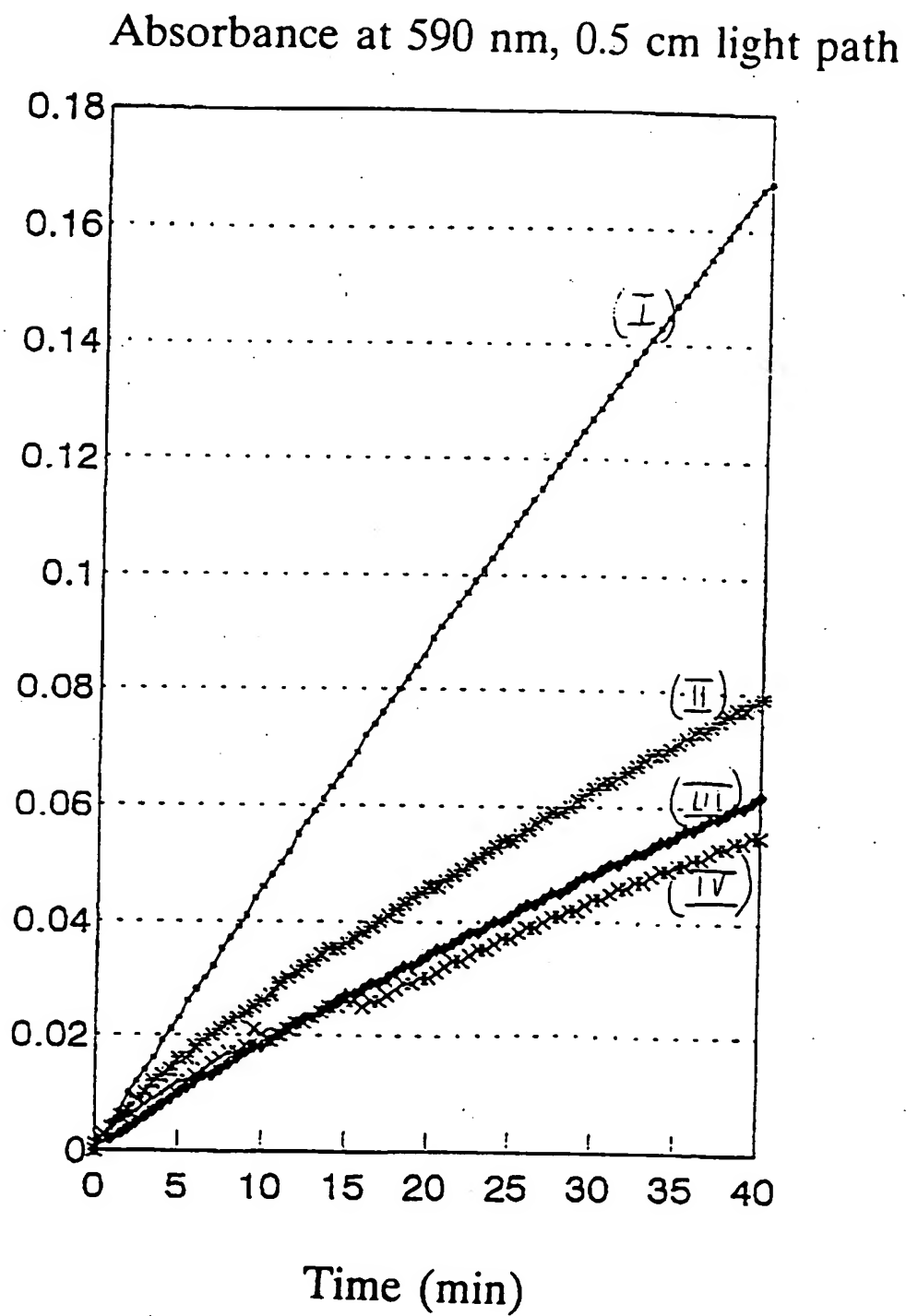


FIG. 1

INTERNATIONAL SEARCH REPORT

International application No.

PCT/DK 95/00344

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: C12N 9/02, C11D 3/386 // C12N 9/02, C12R 1:645

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: C11D, C12N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WPI, CA, BIOSIS, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9313193 A1 (NOVO NORDISK A/S), 8 July 1993 (08.07.93), page 6, line 10 - page 8, line 14, claims 1,9,22 --	1-27
X	WO 9105839 A1 (NOVO NORDISK A/S), 2 May 1991 (02.05.91), page 6, line 6 - line 24; page 7, line 16 - line 21, claims 5,22 --	1-27
A	Patent Abstracts of Japan, Vol 9, No 324, C-320, abstract of JP, A, 60-156385 (Kyowa Hakko Kogyo K.K.), 16 August 1985 (16.08.85) --	1-23

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

& document member of the same patent family

Date of the actual completion of the international search

28 November 1995

Date of mailing of the international search report

30.11.95

Name and mailing address of the ISA/

Swedish Patent Office

Box 5055, S-102 42 STOCKHOLM

Facsimile No. +46 8 666 02 86

Authorized officer

Yvonne Siösteen

Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/DK 95/00344

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 9218683 A1 (NOVO NORDISK A/S), 29 October 1992 (29.10.92), page 3, line 7 - line 20; page 5, line 23 - line 26, and the claims --	1-23
A	WO 9218687 A1 (NOVO NORDISK A/S), 29 October 1992 (29.10.92), page 5, line 8; page 5, line 20 - line 21; page 7, line 6 - line 8 -- -----	1-23

INTERNATIONAL SEARCH REPORT
Information on patent family members

30/10/95

International application No.

PCT/DK 95/00344

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO-A1-	9313193	08/07/93	EP-A-	0617734	05/10/94
			JP-T-	7504694	25/05/95

WO-A1-	9105839	02/05/91	AT-T-	108484	15/07/94
			AU-B-	646645	03/03/94
			AU-A-	6515790	16/05/91
			AU-A-	6516090	16/05/91
			CA-A-	2067748	14/04/91
			CN-A-	1051600	22/05/91
			DE-D,T-	69010691	16/03/95
			EP-A,B-	0495836	29/07/92
			SE-T3-	0495836	
			EP-A,B-	0497794	12/08/92
			SE-T3-	0497794	
			ES-T-	2057593	16/10/94
			JP-T-	5500899	25/02/93
			JP-T-	5503542	10/06/93
			TR-A-	26687	15/05/95
			US-A-	5273896	28/12/93
			WO-A-	9105858	02/05/91

WO-A1-	9218683	29/10/92	NONE		

WO-A1-	9218687	29/10/92	EP-A-	0580707	02/02/94
			JP-T-	6506731	28/07/94
			US-A-	5356437	18/10/94
